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PRIORITY DATE CLAIMED
10 MAY 1999

TITLE OF INVENTION

PROCESS FOR PREPARATION OF PHENOL-MODIFIED ROSIN ESTER, PHENOL-MODIFIED ROSIN ESTER AND USE THEREOF

APPLICANT(S) FOR DO/EO/US

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Applicant herewith submits to the United States Designated /Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(I).
4. ☐ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☐ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☒ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☐ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☐ have not been made and will not be made.
8. ☒ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern other document(s) or information included:

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☒ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☐ A FIRST preliminary amendment.
14. ☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information:

Verification of Translation

(including 32 pages specification, 4 pages claims and 1 page abstract)

First page of WO00/68289

PCT/IB/308

International Search Report

PCT Request

INTERNATIONAL APPLICATION NO.
PCT/JP00/097719359INTERNATIONAL FILING DATE
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17. [X] The following fees are submitted:

Basic National Fee (37 CFR 1.492(a)(1)-(5):

Neither international preliminary examination fee (37 CFR 1.482)

Nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search

Report not prepared by the EPO or JPO (1.492(a)(3)) \$1,000.00

International preliminary examination fee (37 CFR 1.482) not paid to USPTO but

International Search Report prepared by the EPO or JPO (1.492(a)(5)) \$860.00

International preliminary examination fee (37 CFR 1.482) not paid to USPTO but

International search fee (37 CFR 1.445(a)(2)) paid to USPTO (1.492(a)(2)) \$710.00

International preliminary examination fee paid to USPTO (37 CFR 1.482) but all claims did

not satisfy provisions of PCT Article 33(1)-(4) (1.492(a)(1)) \$690.00

International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims

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Claims**Number
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Total Claims 17 -20=

X \$ 18.00

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Independent Claims 1 -3=

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Multiple dependent claim(s) (if applicable)

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a. [X] A check in the amount of \$860 & 40 to cover the above fees is enclosed.

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Deposit Account No. 02-4377. A copy of this sheet is enclosed.**NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or
(b)) must be filed and granted to restore the application to pending status.**

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December 11, 2000

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19,498

Registration No.

-1-

DESCRIPTION

PROCESS FOR PREPARATION OF PHENOL-MODIFIED
ROSIN ESTER, PHENOL-MODIFIED
ROSIN ESTER AND USE THEREOF

5

Field of the Invention

The present invention relates to a novel process
for preparing a phenol-modified rosin ester, a phenol-
modified rosin ester prepared by the process and use
10 thereof.

Background Art

Phenol-modified rosin esters are suitably used
as a binder for printing inks.

15 Heretofore phenol-modified rosin esters have
been prepared by carrying out a procedure for modifying
rosin with phenol and a procedure for esterifying the
rosin. Usually these procedures are conducted by (1) a
process comprising modifying rosin with phenol and then
20 esterifying the rosin with polyhydric alcohol or (2) a
process comprising esterifying rosin with polyhydric
alcohol and then modifying the rosin with phenol. The
phenol-modifying procedure in the processes (1) and (2) is
conducted by (a) a method comprising mixing a solution of
25 resol phenol resin in an organic solvent with rosin or a

polyhydric alcohol ester of rosin with heating at 220 to 270°C to modify the rosin with the phenol, and (b) a method comprising adding phenol and paraformaldehyde to fused rosin or polyhydric alcohol ester of rosin at 80 to 120°C, maintaining the mixture at the same temperature for a specific period in the presence of a catalyst for conversion to resol and heating the mixture to 200 to 250°C to modify the rosin with the phenol.

In the case of the method (a), however, a large amount of calcium hydroxide or like non-volatile metal compound is usually used as the catalyst for conversion to resol in preparing a resol phenol resin as the raw material so that a substantial amount of said compound remains in the phenol-modified rosin ester eventually obtained. Thus, printing inks containing the obtained rosin ester are unsatisfactory in properties, and there is a need for removing the remaining compound from the obtained resol phenol resin to prevent the printing ink from deterioration of properties. For this purpose, a water-washing procedure is indispensable. Therefore, it is essential to use an organic solvent in preparing a resol phenol resin in order to reduce the viscosity of the reaction mixture containing the resol phenol resin and to easily separate and remove the water used for washing.

A solution of resol phenol resin in an organic

solvent is used in the phenol-modifying procedure by the method (a). Thus, when a large amount of the organic solvent solution of the resol phenol resin is supplied to the reaction system at one time, there is a risk of

5 causing bump of organic solvent, and the temperature of reaction system may be lowered due to a high latent heat in vaporization of organic solvent. In view of this possibility, the resin solution should be added by degrees to the reaction system, involving not only a drawback of a

10 prolonged time period taken in preparation of the resin, but also disadvantages of using an organic solvent in preparing a resol phenol resin and necessitating additional procedures and increased equipment costs for washing with water and for disposal of waste water.

15 On the other hand, in the case of the method (b), a phenol resin is made into resol concurrently with modification in the phenol-modifying procedure. Consequently the method (b) is free from disadvantages of additional procedures and increased equipment costs as in

20 the method (a) and is economical. However, because of self-condensation of phenol resin and addition reaction of superfluous formalin with the rosin, the method (b) entails difficulty in obtaining a phenol-modified rosin ester having a high molecular weight and a high solubility

25 among the properties required these days by binders for

printing inks.

Disclosure of the Invention

An object of the present invention is to provide
5 a novel process for preparing a phenol-modified rosin
ester which has overcome the above-mentioned prior art
problems.

Another object of the present invention is to
provide a phenol-modified rosin ester prepared by said
10 process.

A further object of the present invention is to
provide a binder for printing inks which contains said
phenol-modified rosin ester and a printing ink containing
the binder for printing inks.

Other objects and features of the present
15 invention will become apparent from the following
description.

According to the present invention, there is
provided a process for preparing a phenol-modified rosin
20 ester, the process comprising the step of reacting rosin
with phenol, formaldehyde and polyhydric alcohol, the
process being characterized in that rosin or a polyhydric
alcohol ester of rosin is reacted with a resol phenol
resin prepared in the presence of a volatile base catalyst.

25 According to the present invention, there is

provided a phenol-modified rosin ester prepared by said process.

According to the present invention, there are provided a binder for printing inks which binder contains
5 said phenol-modified rosin ester and a printing ink containing said binder for printing inks.

The present inventor conducted extensive research to overcome the foregoing problems, directing attention to resol phenol resins useful as the raw
10 material for a phenol-modified rosin ester and found the following. When the above-specified resol phenol resin is used in the phenol-modified procedure by the method (a), a high molecular weight phenol-modified rosin ester having a high solubility in a solvent for inks can be prepared in a
15 short time using relatively simple equipment. Furthermore, the obtained phenol-modified rosin ester is suitable as a binder for printing inks which binder has improved printability.

The present invention was completed based on
20 these novel findings.

According to the process of the invention for preparing a phenol-modified rosin ester, it is essential in the phenol-modifying procedure to use a resol phenol resin which is prepared by reacting phenol with
25 formaldehyde in the presence of a volatile base serving as

a catalyst, optionally in a closed reactor under an increased pressure. The resol phenol resin used gives the following various advantages.

The foregoing resol phenol resin is free of the

5 non-volatile metal compound conventionally used as a catalyst for conversion to resol, so that the resol phenol resin need not be washed with water after preparation of the resol phenol resin. Consequently the resol phenol resin can be prepared without use of a solvent. When the

10 resol phenol resin is prepared in a closed reactor under an increased pressure, the formaldehyde and volatile base catalyst can be prevented from dissipating away from the reaction system and the conversion to resol can be conducted in a very short time. Further, the obtained

15 resol phenol resin is substantially free of a solvent so that the resin can be added at a unlimited speed in the phenol-modifying procedure, whereby the phenol-modifying time can be pronouncedly shortened.

The phenol-modified rosin ester prepared using

20 said resol phenol resin as the raw material can unexpectedly exhibit excellent properties for printability when used as a binder for printing inks, as compared with conventional phenol-modified rosin esters, presumably because of the foregoing advantages in the production and

25 the residual volatile base catalyst.

In the present invention, rosin or a polyhydric alcohol ester of rosin is reacted with a specific resol phenol resin to give a phenol-modified rosin ester.

Rosins useful herein as the raw material for the
5 phenol-modified rosin ester are various and include, for example, gum rosin, wood rosin, tall oil rosin, distilled rosin, hydrogenated rosin, polymerized rosin, disproportionated rosin, etc.

Examples of phenols useful herein as the raw
10 material for the resol phenol resin are phenol, p-octylphenol, p-nonylphenol, p-dodecylphenol, p-butylphenol, bisphenol A, phenylphenol, p-aminophenol, p-ethylphenol, etc.

Formaldehydes useful herein as the raw material
15 for the resol phenol resin include, for example, paraformaldehyde and an aqueous solution of formalin. The amount of the formaldehyde used is about 0.5 to about 3 moles (calculated as formaldehyde in the case of paraformaldehyde) per mole of phenol. When the amount of
20 the formaldehyde used is less than 0.5 mole, the obtained phenol-modified rosin ester is rendered more soluble but is likely to lower in the molecular weight because the rosin ester contains considerable amounts of unreacted phenol and rosin residues. When the amount exceeds 3
25 moles, the obtained phenol-modified rosin ester has a high

molecular weight but is likely to lower in the solubility. From these viewpoints, preferably the amount of formaldehyde ranges from the upper limit of not higher than 2.5 moles to the lower limit of not lower than 1.5
5 moles.

In the present invention, it is essential to use the resol phenol resin prepared by reacting phenol with formaldehyde in the presence of a volatile base catalyst.

There is no limitation on other reaction
10 conditions for preparing the resol phenol resin than use of a volatile base as the catalyst. A preferred reaction temperature for conversion to resol is about 80 to about 200°C. At lower than 80°C, the reaction for conversion to resol proceeds at an exceedingly low rate, whereas at
15 higher than 200°C, the reaction becomes difficult to control. From these viewpoints, a more preferred reaction temperature for conversion to resol is 160°C in the upper limit and 90°C in the lower limit.

When the reaction for conversion to resol is
20 carried out in said reaction temperature range, usually the formaldehyde is vaporized and the water generated in the reaction is boiled, so that a pressure-resistant reactor is preferably used. Further, if the reaction is conducted in a closed reactor under an increased pressure,
25 it is possible to prevent dissipation of phenol, formalin

and volatile base catalyst from the reaction system. Thus the reaction time can be shortened by elevating the reaction temperature. For example, under an increased pressure of 1.0 MPa, the reaction can be performed at about 180°C. For example, the reaction conventionally taking a few hours can be completed only in 5 minutes. Moreover, since the viscosity of the resin is lowered at a high temperature, the reaction for conversion to resol can be suitably conducted without use of a solvent. In this case, the yield is further increased and the procedure is made more economical because the solvent inherently unwanted need not be used. However, a solvent is allowable to use in the reaction for conversion to resol and a solvent may be suitably used, if necessary.

15 In the present invention, if the resol phenol resin to be used as the raw material has too low a molecular weight, the rosin ester prepared using the resol phenol resin is given a low solubility. If the resol phenol resin has too high a molecular weight, the rosin ester is given a low viscosity. Consequently, a resol phenol resin of about 500 to about 2,500 in molecular weight is preferable to use.

 The volatile base catalyst for use in preparing a resol phenol resin is preferably one having a boiling point or a decomposition temperature in the range of

usually not higher than about 200°C. Specific examples of the volatile base catalyst are an aqueous solution of ammonia, hexamethylenetetramine and like ammonia-generating substances; methylamine, ethylamine, 5 propylamine, butylamine, amylamine, hexylamine, heptylamine, octylamine, nonylamine, decylamine, undecylamine, dodecylamine, tridecylamine, tetradecylamine, pentadecylamine, cetylamine and like aliphatic primary amines; dimethylamine, diethylamine, dipropylamine, 10 diisopropylamine, dibutylamine, diamylamine and like aliphatic secondary amines; trimethylamine, triethylamine and like aliphatic tertiary amines; allylamine, diallylamine, triallylamine and like aliphatic unsaturated amines; cyclopropylamine, cyclobutylamine, 15 cyclopentylamine, cyclohexylamine and like alicyclic amines; benzylamine, dibenzylamine, tribenzylamine, diphenylamine, triphenylamine, α -naphthylamine, β -naphthylamine and like aromatic amines; monoethanolamine, dimethylaminoethanol, diethanolamine, triethanolamine and 20 like amino alcohols; etc. Also usable are, for example, ethylenediamine and like polyvalent amines; pyridine, piperidine, piperazine, pyrazine, aniline and so on. Further usable are salts of the above-exemplified compounds with weak acids and quaternary ammonium salts of 25 the above-exemplified compounds. Among them, preferred

are trimethylamine, diethylamine, triethylamine, monoethanolamine, dimethylaminoethanol, diethanolamine and the like from the viewpoints of catalytic activity, boiling point, safety and handleability.

5 The amount of the volatile base catalyst used is suitably determinable depending on the time involved in preparing the resol phenol resin, and the nitrogen residue content as described later among the properties of the obtained phenol-modified rosin ester.

10 The amount of the resol phenol resin used is usually at least 20 parts by weight per 100 parts of rosin or polyhydric alcohol ester of rosin. If less than 20 parts by weight of the resin is used, it would be difficult to obtain the contemplated high molecular weight
15 rosin ester. On the other hand, the upper limit in the amount of resol phenol resin is usually not higher than 200 parts by weight per 100 parts of rosin or polyhydric alcohol ester of rosin from the standpoint of costs.

 Examples of polyhydric alcohols useful as the
20 component for esterification of rosin are glycerine, diglycerine, polyglycerine, pentaerythritol, dipentaerythritol, diethylene glycol, ethylene glycol, trimethylolethane, trimethylolpropane, etc. The amount of the polyhydric alcohol used is about 0.3 to about 1.5 in
25 terms of the equivalent ratio (-OH equivalent of

polyhydric alcohol/-COOH equivalent of rosin). The equivalent ratio of less than 0.3 makes it difficult to obtain the contemplated high molecular weight rosin ester, whereas more than 1.5 tends to impart a lower solubility to the obtained phenol-modified rosin ester. In view of the above, a preferred amount of the polyhydric alcohol used is about 0.5 to about 1.3 in terms of said equivalent ratio.

Examples of components which are optionally usable in preparing the phenol-modified rosin ester of the invention are petroleum resins such as those obtained from dicyclopentadiene or the like; linseed oil, soybean oil, tung oil, castor oil, rapeseed oil, cottonseed oil, coconut oil and like oils; oleic acid, stearic acid, lauric acid, isostearic acid, dimer acid and like fatty acids; etc. The amount of these optional components used is not higher than about 100 parts by weight per 100 parts by weight of rosin.

According to the present invention, the phenol-modified rosin ester is prepared by reacting rosin with a resol phenol resin and polyhydric alcohol. The process for preparing the rosin ester can be properly selected from, for example, (1) a process comprising the steps of modifying rosin with a resol phenol resin and then esterifying the modified rosin with polyhydric alcohol,

(2) a process comprising the steps of esterifying rosin with polyhydric alcohol to give a polyhydric alcohol ester of rosin, and then modifying the ester with a resol phenol resin, and (3) a process comprising the steps of charging
5 into a reactor rosin, a resol phenol resin and polyhydric alcohol at the same time and reacting them.

The reaction conditions for preparing the phenol-modified rosin ester are not limited and include conventional reaction conditions. Generally, the reaction
10 temperature for modifying the rosin or polyhydric alcohol ester of rosin with phenol is usually about 180 to about 270°C, and the reaction time is usually about 3 to about 5 hours. The temperature for esterification is usually about 230 to about 280°C and the reaction time is about 10
15 to about 16 hours. In said esterification procedure, an esterification catalyst can be used when so required.

The phenol-modified rosin ester of the present invention used as a binder for printing inks is in no way inferior in properties to the rosin esters prepared by the
20 above-mentioned conventional method (a). According to the process of the invention for preparing the phenol-modified rosin ester, a high molecular weight and a highly soluble phenol-modified rosin ester can be easily prepared.

When the phenol-modified rosin ester is used as
25 a binder for printing inks, the property values of the

phenol-modified rosin ester greatly affect the properties of printing inks such as resistance to emulsification and resistance to smudging on printed paper sheets.

Consequently the property values of the phenol-modified
5 rosin ester should be carefully determined and some property values are desirably in the following ranges.

The phenol-modified rosin ester of the invention has desirably an acid value of 10 to 40 mg KOH/g, preferably 15 to 30 mg KOH/g, a weight average molecular
10 weight of 10,000 to 500,000, preferably 20,000 to 200,000, calibrated with polystyrene standard samples, and a softening point (ring and ball method) of 140 to 190°C, preferably 150 to 180°C.

The phenol-modified rosin ester of the invention
15 has desirably a nitrogen residue content (as measured by microanalysis of total nitrogen by a catalyst oxidation conversion method) in the range of 10 to 1,000 ppm, preferably 30 to 500 ppm. The nitrogen residue content results from the volatile base catalyst used in preparing
20 the resol phenol resin. Therefore, the phenol-modified rosin ester contains the residue in the original form of volatile base catalyst or in any form in which it exists as bonded to atoms in the molecules of the rosin ester.

The phenol-modified rosin ester of the invention
25 has desirably a solubility (25°C) in a petroleum solvent

(boiling point of 276 to 318°C, aniline point 69°C) in the range of at least 2 times, preferably at least 5 times.

Useful petroleum solvents include, for example, "No.5 Solvent" (trade name, manufactured by Nippon Mitsubishi

5 Oil Corporation) which is used as a solvent for inks. The solubility herein referred to is evaluated as follows. A 1.0 g quantity of the phenol-modified rosin ester is charged into a test tube, 2.0 g of No.5 Solvent is added to the test tube and the mixture is heated to 180°C to
10 form a solution, followed by cooling to 25°C. Then, the solvent is further added when required, and the procedure is terminated when the solution has become turbid. The total amount (g) of the solvent per gram of the rosin ester (g/g) upon termination of procedure is indicated.

15 If the property values of the phenol-modified rosin ester are outside the above-specified ranges, the printing ink containing the rosin ester as a binder would exhibit poor printing properties. Hence the rosin ester of properties outside said ranges is undesirable.

20 The phenol-modified rosin ester prepared according to the invention can be suitably used, for example, as a binder for printing inks. The printing ink containing the rosin ester as a binder can be used especially for offset printing and for planography
25 printing (dry), letterpress printing, gravure printing or

the like.

To prepare a binder for printing inks using the phenol-modified rosin ester of the invention, the rosin ester is dissolved in a petroleum solvent, drying oil or
5 semi-drying oil with heating, and optionally a gelling agent is suitably added. In formulating a printing ink using said binder for printing inks, conventional pigments, non-aromatic solvents, drying oils, abrasion resistance improvers, dryers and like additives may be incorporated
10 in the printing ink and milled.

Best Mode for Carrying Out the Invention

The following examples are given to clarify the present invention in more detail. The parts and
15 percentages used in the examples are all by weight.

Comparative Preparation Example 1

(Example wherein a conventional resol phenol resin is prepared without washing with water)

A reactor equipped with a stirrer, a reflux
20 condenser having a water separator and a thermometer was charged with 4,000 parts of p-octylphenol, 1,800 parts of 92% paraformaldehyde and 2,000 parts of xylene. The mixture was heated to 60°C to form a solution. Then 72 parts of calcium hydroxide was added as a catalyst for
25 conversion to resol. Thereafter the mixture was heated to

90°C to undergo a reaction for 4 hours, giving a solution of a resol phenol resin having a weight average molecular weight of 740 (solid content 65%).

Comparative Preparation Example 2

- 5 (Example wherein a conventional resol phenol resin is prepared by washing with water)

A predetermined amount of hydrochloric acid was added to 7,500 parts of the resol phenol resin solution prepared in Comparative Preparation Example 1 to adjust a
10 pH of the contents to about 4. After washing the contents with 3,000 parts of warm water, the contents were left to stand for 2 hours to separate the oil layer, giving a solution of a resol phenol resin having a weight average molecular weight of 745 (solid content 65%).

- 15 Preparation Example 1

(Example wherein a resol phenol resin is prepared under an increased pressure)

A pressure-resistant reactor (1.5 MPa) equipped with a stirrer, an internal pressure meter, a reflux
20 condenser having a water separator and a thermometer was charged with 4,000 parts of p-octylphenol and 1,300 parts of 92% paraformaldehyde. The mixture was heated to 80°C in a closed reactor under an increased pressure to form a solution, and 120 parts of triethylamine was added as a
25 catalyst for conversion to resol. Then the mixture was

heated to 145°C to undergo a reaction for 5 minutes, giving a solution of a resol phenol resin having a weight average molecular weight of 746 (solid content 95%).

Preparation Example 2

- 5 (Example wherein a resol phenol resin is prepared under an increased pressure)

The foregoing pressure-resistant reactor was charged with 4,000 parts of p-octylphenol and 1,050 parts of 92% paraformaldehyde. The mixture was heated to 80°C in
10 a closed reactor under an increased pressure to form a solution, and 120 parts of triethylamine was added as a catalyst for conversion to resol. Then the mixture was heated to 125°C to undergo a reaction for 30 minutes, giving a solution of a resol phenol resin having a weight
15 average molecular weight of 755 (solid content 95%).

Preparation Example 3

(Example wherein a resol phenol resin is prepared under an increased pressure)

The foregoing pressure-resistant reactor was
20 charged with 4,000 parts of p-octylphenol and 1,050 parts of 92% paraformaldehyde. The mixture was heated to 80°C in a closed reactor under an increased pressure to form a solution, and 70 parts of trimethylamine was added as a catalyst for conversion to resol. Then the mixture was
25 heated to 125°C to undergo a reaction for 30 minutes,

giving a solution of a resol phenol resin having a weight average molecular weight of 782 (solid content 95%).

Preparation Example 4

(Example wherein a resol phenol resin is prepared under an
5 atmospheric pressure)

A reactor was charged with 4,000 parts of p-octylphenol and 1,050 parts of 92% paraformaldehyde. The mixture was heated to 80°C to form a solution, and 110
10 parts of dimethylaminoethanol was added as a catalyst for conversion to resol. Then the mixture was heated to 90°C to undergo a reaction for 4 hours, giving a solution of a resol phenol resin having a weight average molecular weight of 715 (solid content 95%).

Table 1 shows the weight average molecular
15 weight of the resol phenol resins prepared in the above-described Preparation Examples, and the reaction time and the reaction temperature required in preparing the resol phenol resins.

Table 1

	Weight average molecular weight	Reaction time (minute)	Reaction temperature (°C)
Prep.Ex. 1	746	5	145
Prep.Ex. 2	755	30	125
Prep.Ex. 3	782	30	125
Prep.Ex. 4	715	240	90
Comp.Prep.Ex. 1	740	240	90
Comp.Prep.Ex. 2	745	240	90

Example 1

1,800 parts of gum rosin was charged into a
5 reactor equipped with a stirrer, a reflux condenser having
a water separator and a thermometer and was fused with
heating to 230°C in a nitrogen atmosphere. Then 170 parts
of pentaerythritol was added and homogeneously mixed, and
4 parts of calcium hydroxide was added as an
10 esterification catalyst. Thereafter the mixture was
heated to 280°C. The water produced was collected by the
reflux condenser having a water separator. A reaction was
conducted at the same temperature for 8 hours and the
reaction mixture was cooled to 250°C when the acid value
15 reached not higher than 25 mg KOH/g. Subsequently 1,100
parts of the resol phenol resin solution prepared in
Preparation Example 1 (solid content 95%) was added
dropwise over a period of 4 hours at the same temperature.
The mixture was held at the same temperature awhile.

After it was confirmed that a predetermined resin viscosity was attained, a phenol-modified rosin ester was produced. The obtained rosin ester was found to have an acid value of 22 mg KOH/g, a weight average molecular weight of 43,000 (calibrated with polystyrene standard samples), a softening point of 169°C (ring and ball method), a nitrogen residue content of 200 ppm (measured by microanalysis of total nitrogen by catalyst oxidation conversion method, a measuring device used: "TN-10 Model" manufactured by Mitsubishi Chemical Corp.) and a solubility (25°C) of not less than 20 times when using "No.5 Solvent" (trade name, product of Nippon Mitsubishi Oil Corporation) as a petroleum hydrocarbon solvent (boiling point range 276 to 313°C, aniline point 69°C).

15 Example 2

1,800 parts of gum rosin was charged into the foregoing reactor and was fused with heating to 250°C in a nitrogen atmosphere. Subsequently 1,050 parts of the resol phenol resin solution prepared in Preparation

20 Example 2 (solid content 95%) was added dropwise over a period of 3 hours. Then 170 parts of glycerine was added and the mixture was held at 250°C. The water produced was collected by the reflux condenser having a water separator. A reaction was conducted at the same temperature for 8

25 hours. After it was confirmed that the acid value reached

not higher than 25 mg KOH/g, a phenol-modified rosin ester was obtained.

Example 3

1,800 parts of gum rosin was charged into the foregoing reactor and was fused with heating to 250°C in a nitrogen atmosphere. Subsequently 1,050 parts of the resol phenol resin solution prepared in Preparation Example 3 (solid content 95%) was added dropwise over a period of 3 hours. Then 170 parts of glycerine was added and the mixture was held at 250°C. The water produced was collected by the reflux condenser having a water separator. A reaction was conducted at the same temperature for 8 hours. After it was confirmed that the acid value reached not higher than 25 mg KOH/g, a phenol-modified rosin ester was obtained.

Example 4

1,800 parts of gum rosin was charged into the foregoing reactor and was fused with heating to 250°C in a nitrogen atmosphere. Subsequently 1,050 parts of the resol phenol resin solution prepared in Preparation Example 4 (solid content 95%) was added dropwise over a period of 3 hours. Then 170 parts of glycerine was added and the mixture was held at 250°C. The water produced was collected by the reflux condenser having a water separator. A reaction was conducted at the same temperature for 8

hours. After it was confirmed that the acid value reached not higher than 25 mg KOH/g, a phenol-modified rosin ester was obtained.

Example 5

5 1,800 parts of gum rosin was charged into the foregoing reactor and was fused with heating to 230°C in a nitrogen atmosphere. Subsequently 85 parts of pentaerythritol and 10 parts of glycerine were added and homogeneously mixed. The mixture was heated to 285°C. The
10 water produced was collected by the reflux condenser having a water separator. A reaction was conducted at the same temperature for 3 hours. When the acid value reached 80 mg KOH/g, the mixture was cooled to 250°C. Subsequently 1,200 parts of the resol phenol resin solution prepared in
15 Preparation Example 2 (solid content 95%) was added dropwise over a period of 4 hours. Then 80 parts of glycerine was added and the reaction was continued for about 6 hours. The water produced was collected by the reflux condenser having a water separator. After it was
20 confirmed that the acid value reached not higher than 20 mg KOH/g, a phenol-modified rosin ester was obtained.

Comparative Example 1

 1,800 parts of gum rosin was charged into the foregoing reactor and was fused with heating to 230°C in a
25 nitrogen atmosphere. Subsequently 170 parts of

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solution prepared in Comparative Preparation Example 2 (solid content 65%) was added dropwise over a period of 12 hours. The xylene and water were collected by the reflux condenser having a water separator. On completion of

5 dropwise addition of resol phenol resin solution, the mixture was held at the same temperature for 1 hour after which a phenol-modified rosin ester was obtained.

Comparative Example 3

1,800 parts of gum rosin was charged into the

10 foregoing reactor and was fused with heating to 250°C in a nitrogen atmosphere. Subsequently 1,530 parts of the resol phenol resin solution prepared in Comparative Preparation Example 2 (solid content 65%) was added dropwise over a period of 7 hours. Then, 170 parts of

15 glycerine was added and a reaction was conducted at the same temperature for 12 hours, giving a phenol-modified rosin ester. The xylene and water were collected by the reflux condenser having a water separator. The rosin ester was found to have an acid value of 24 mg KOH/g, a

20 softening point of 175°C, a solubility (25°C) of 11.5 times in "No.5 Solvent" described above, and a weight average molecular weight of 65,000.

Comparative Example 4

1,800 parts of gum rosin was charged into the

25 foregoing reactor and was fused with heating to 230°C in a

nitrogen atmosphere. Subsequently 85 parts of pentaerythritol and 10 parts of glycerine were added and homogeneously mixed. The mixture was heated to 285°C. The water produced was collected by the reflux condenser

5 having a water separator. A reaction was conducted at the same temperature for 3 hours. When the acid value reached 80 mg KOH/g, the mixture was cooled to 250°C. Subsequently 1,600 parts of the resol phenol resin solution prepared in Comparative Preparation Example 2 (solid content 65%) was

10 added dropwise over a period of 7 hours, followed by addition of 80 g of glycerine. The reaction was continued for about 6 hours. The water produced was collected by the reflux condenser having a water separator. When the acid value reached not higher than 20 mg KOH/g, the reflux

15 was terminated, and the mixture was held at the same temperature awhile. After it was confirmed that the predetermined resin viscosity was attained, a phenol-modified rosin ester was obtained.

Table 2 shows some properties of the phenol-

20 modified rosin esters obtained in the Examples and Comparative Examples and the time taken in preparing the rosin ester.

Table 2

	Acid value (mg KOH/g)	Softening point(°C)	Weight average Molecular weight
Ex. 1	22	169	43,000
Ex. 2	24	175	65,000
Ex. 3	24	174	60,000
Ex. 4	24	175	62,000
Ex. 5	19	176	122,000
Comp.Ex. 1	22	170	44,000
Comp.Ex. 2	22	172	43,000
Comp.Ex. 3	24	175	65,000
Comp.Ex. 4	19	178	120,000

Table 2 (continued)

	Nitrogen residue content (ppm)	Solubility (g/g)	Production time(hr)
Ex. 1	200	20<	18
Ex. 2	120	12.2	16
Ex. 3	100	14	16
Ex. 4	150	12.0	16
Ex. 5	160	20<	20
Comp.Ex. 1	-	18	24
Comp.Ex. 2	-	20<	24
Comp.Ex. 3	-	11.5	24
Comp.Ex. 4	-	20<	27

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Preparation of Printing Ink

A varnish was prepared by mixing 43 parts of each of the phenol-modified rosin esters prepared in Examples 1 to 5 and Comparative Examples 1 to 4 with 20

parts of linseed oil and 45 parts of "AF Solvent No.6"
(trade name for a product of Nippon Mitsubishi Oil
Corporation, hydrogenated petroleum hydrocarbon solvent
having a boiling point of 302 to 321°C) at 180°C to form a
5 solution. Then, 0.5 part of ethylacetoacetate aluminum
diisopropylate (trade name "ALCH", product of Kawaken Fine
Chemicals Co., Ltd.) was added to 100 parts of the varnish.
The mixture was reacted at 190°C for 1 hour to give a
varnish gel. A printing ink was prepared by milling the
10 varnish gel with a three-roll mill using the following
components in the proportions shown below:

	Crimson pigment	18 parts
	Foregoing varnish gel	70-78 parts
	"AF Solvent No.6"	3-11 parts
15	Drier	1 part

The ink thus obtained was suitably adjusted to a
tack value of 8.5 ± 0.5 and a flow value of 18 ± 0.5 .

"Brilliant Carmine 6B" (trade name, a product of
Toyo Ink Manufacturing Co., Ltd.) was used as the crimson
20 pigment and cobalt naphthenate was used as the drier.

The obtained inks were tested for performance by
the following methods.

Gloss: The ink (0.27 ml) was applied to the surface of a
sheet of wood-free paper using "RI Tester" (trade name,
25 product of Ishikawajima Sangyo Kikai Co., Ltd.) according

- to JIS K 5701. The obtained inked sheet sample was dried in the atmosphere at 25°C for 24 hours, and 60-degree specular gloss (%) was measured according to JIS Z 8741.
- Resistance to emulsification: The ink (3.9 ml) was applied
- 5 to a kinetic emulsification tester (product of Nihon Rheology Kiki Co., Ltd.). Pure water was supplied at 200 r.p.m. and at a rate of 5 ml/min. The water content in the ink was measured by an infrared moisture meter. The emulsification ratio was calculated by the following
- 10 equation.
- Emulsification ratio (%) = $\frac{\text{weight of water in ink}}{\text{weight of water in ink} + \text{weight of ink}} \times 100$
- Resistance to misting: In an air-conditioned chamber adjusted to 25°C was the ink (2.7 ml) applied to a roll of
- 15 the inkometer as defined in JIS K 5701, and the roll was manually turned to spread the ink uniformly over the roll. A blank sheet of paper was held at a position away by 10 cm from the roll while the roll was rotated at 1,200 r.p.m. for 2 minutes. The amount of ink scattered over the blank
- 20 sheet of paper was visually evaluated according to the following criteria. A: Substantially no ink was scattered, B: A substantial amount of ink was scattered. A preferred ink is one scarcely scattered as is the case with the ink used in a printing machine actually operated.
- 25 Dryness degree: The ink (0.27 ml) was applied to the

surface of a sheet of sulfate paper using "RI Tester"
(trade name, product of Ishikawajima Sangyo Kikai Co.,
Ltd.) according to JIS K 5701. Another sheet of sulfate
paper was superposed on the inked surface of sulfate paper
5 sheet and was wound on a rotary drum of a C-type dryness
tester (product of Toyoseiki Seisakusho Co., Ltd.) in such
a manner that the paper sheet superposed on the inked
paper sheet externally existed over the drum. A weight
and a compression gear were slowly placed on the external
10 sheet, and then the drum was rotated. The drying time
(hr) is equal to a period of time taken until
substantially no trace of the gear is left on the external
sheet.

Resistance to smudging on printed paper sheets: Using a
15 light offset printing press (product of Ryobi Ltd., trade
name "2800 CD"), 1,000 paper sheets were printed while the
water-supplying dial was fully throttled until immediately
before the printed sheet became smudged. The degree of
smudging on printed paper sheets was visually evaluated
20 according to the following criteria. A: No smudging; B:
Slightly smudged; C: Markedly smudged.

Table 3 shows the results of performance tests.

Table 3

	Gloss (%)	Emulsification ratio (%)	Resistance to misting	Drying time (hr)	Resistance to smudging on printed paper sheets
Ex. 1	70	40	B	5	A
Ex. 2	60	38	A	4.5	A
Ex. 3	61	40	A	4.5	A
Ex. 4	60	42	A	4.5	A
Ex. 5	75	30	A	4.5	A
Comp.Ex.1	35	65	A	4.5	C
Comp.Ex.2	62	52	B	5	A
Comp.Ex.3	58	40	A	4.5	B
Comp.Ex.4	70	35	A	4.5	A

The phenol-modified rosin ester of Comparative Example 1 was prepared using the resol phenol resin of Comparative Preparation Example 1 (without washing with water) obtained in the presence of the conventional metal compound catalyst. As apparent from Table 3, a printing ink containing this rosin ester as a binder was inferior in properties such as gloss, resistance to emulsification (emulsification ratio) and resistance to smudging on printed paper sheets. Further, the phenol-modified rosin esters of Comparative Examples 2 to 4 were prepared using the resol phenol resin of Comparative Preparation Example 2 (involving washing with water) obtained in the presence of the conventional metal compound as a catalyst. Printing inks containing the rosin esters as a binder were

not fully satisfactory in printing properties. On the other hand, when the phenol-modified rosin esters of Examples 1 to 5 according to the invention were used, evidently the obtained printing inks containing these
5 rosin esters as a binder were remarkably superior in printing properties.

According to the process of the present invention, the resol phenol resin useful as the raw material can be prepared in an exceedingly shorter time,
10 and the reaction for conversion to resol can be carried out without use of a solvent. According to the process of the present invention, the equipment for preparing a phenol-modified rosin ester can be simplified and the rosin ester can be prepared in a markedly short time. The
15 phenol-modified rosin ester of the present invention has the same constants for resins as the phenol-modified rosin esters prepared by conventional processes, but can exhibit excellent properties when used as a binder for printing inks.

CLAIM

1. A process for preparing a phenol-modified
rosin ester, the process comprising the step of reacting
rosin with phenol, formaldehyde and polyhydric alcohol,
5 the process being characterized in that rosin or a
polyhydric alcohol ester of rosin is reacted with a resol
phenol resin prepared in the presence of a volatile base
catalyst.
- 10 2. The process according to claim 1, wherein the
resol phenol resin is one prepared by reacting phenol with
formaldehyde at 80 to 200°C.
3. The process according to claim 1, wherein the
15 resol phenol resin is one prepared by reacting phenol with
formaldehyde in a closed reactor under an increased
pressure.
4. The process according to claim 1, wherein the
20 volatile base catalyst in the resol phenol resin is amine.
5. The process according to claim 1, wherein the
resol phenol resin is one prepared by reacting 1 mole of
phenol with 0.5 to 3 moles of formaldehyde.

6. The process according to claim 1, wherein the
rosin is at least one species selected from the group
consisting of gum rosin, wood rosin, tall oil rosin,
distilled rosin, hydrogenated rosin, polymerized rosin and
5 disproportionated rosin.

7. The process according to claim 1, wherein the
phenol-modified rosin ester is one prepared by reacting
rosin with a resol phenol resin, and reacting the reaction
10 mixture with polyhydric alcohol.

8. The process according to claim 1, wherein the
phenol-modified rosin ester is one prepared by reacting
rosin with polyhydric alcohol to give a polyhydric alcohol
15 ester of rosin, and reacting the same with a resol phenol
resin.

9. The process according to claim 1, wherein the
rosin or polyhydric alcohol ester of rosin is reacted with
20 the resol phenol resin at 180 to 270°C.

10. A phenol-modified rosin ester prepared by
the process as defined in claim 1.

25 11. The phenol-modified rosin ester according to

claim 10 which has an acid value of 10 to 40 mg KOH/g.

12. The phenol-modified rosin ester according to claim 10 which has a weight average molecular weight of 10,000 to 500,000 as calibrated with polystyrene standard samples.

13. The phenol-modified rosin ester according to claim 10 which has a softening point (ring and ball method) of 140 to 190°C.

14. The phenol-modified rosin ester according to claim 10, wherein the nitrogen residue content resulting from the volatile base catalyst (measured by microanalysis of total nitrogen by a catalyst oxidation conversion method) is 10 to 1,000 ppm.

15. The phenol-modified rosin ester according to claim 10 which has a solubility in a petroleum hydrocarbon solvent (boiling point 276 to 313°C, aniline point 69°C) in the range of at least 2 times.

16. A binder for printing inks which contains the phenol-modified rosin ester of claim 10.

17. A printing ink containing the binder for
printing inks as defined in claim 16.

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ABSTRACT

This invention provides a process for preparing a phenol-modified rosin ester, the process comprising the step of reacting rosin with phenol, formaldehyde and
5 polyhydric alcohol, the process being characterized in that rosin or a polyhydric alcohol ester of rosin is reacted with a resol phenol resin prepared in the presence of a volatile base catalyst, the invention further provides a phenol-modified rosin ester prepared by the
10 above-mentioned process, and the invention also provides a binder for printing inks which contains the phenol-modified rosin ester and further provides a printing ink containing the binder for printing inks.

**COMBINED DECLARATION
AND POWER OF ATTORNEY****(Original, Design, National Stage of PCT, Divisional, Continuation or C-I-P Application)**

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name; I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

PROCESS FOR PREPARATION OF PHENOL-MODIFIED ROSIN ESTER, PHENOL-MODIFIED ROSIN ESTER AND
This declaration is of the following type: USE THEREOF

- ☐ original
☐ design
☒ national stage of PCT.
☐ divisional
☐ continuation
☐ continuation-in-part (C-I-P)

the specification of which: *(complete (a), (b), or (c))*

(a) ☐ is attached hereto.

(b) ☐ was filed on as Application Serial No. and was amended on *(if applicable)*.

(c) ☒ was described and claimed in PCT International Application No. filed on and was amended on
(if applicable). PCT/JP00/02918 May 2, 2000

Acknowledgement of Review of Papers and Duty of Candor

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the patentability of the subject matter claimed in this application in accordance with Title 37, Code of Federal Regulations § 1.56.

☐ In compliance with this duty there is attached an information disclosure statement. 37 CFR 1.98.

Priority Claim

I hereby claim foreign priority benefits under Title 35, United States Code, § 119(a)-(d) of any foreign application(s) for patent or inventor's certificate or of any PCT International Application(s) designating at least one country other than the United States of America listed below and have also identified below any foreign application(s) for patent or inventor's certificate or any PCT International Application(s) designating at least one country other than the United States of America filed by me on the same subject matter having a filing date before that of the application on which priority is claimed

(complete (d) or (e))

(d) ☐ no such applications have been filed.

(e) ☒ such applications have been filed as follows:

BAKER & BOTTS, L.L.P.
FILE NO.: RBH

PRIOR FOREIGN/PCT APPLICATION(S) FILED WITHIN 12 MONTHS (6 MONTHS FOR DESIGN) PRIOR TO SAID APPLICATION				
COUNTRY	APPLICATION NO.	DATE OF FILING (day, month, year)	DATE OF ISSUE (day, month, year)	PRIORITY CLAIMED UNDER 35 USC 119
Japan	1999-127983	10/05/1999		<input checked="" type="checkbox"/> YES NO <input type="checkbox"/>
				<input type="checkbox"/> YES NO <input type="checkbox"/>
				<input type="checkbox"/> YES NO <input type="checkbox"/>
ALL FOREIGN APPLICATION(S), IF ANY, FILED MORE THAN 12 MONTHS (6 MONTHS FOR DESIGN) PRIOR TO SAID APPLICATION				
				<input type="checkbox"/> YES NO <input type="checkbox"/>
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Claim for Benefit of Prior U.S. Provisional Application(s)

I hereby claim the benefit under Title 35, United States Code, § 119(e) of any United States provisional application(s) listed below:

Provisional Application Number	Filing Date

Claim for Benefit of Earlier U.S./PCT Application(s) under 35 U.S.C. 120

(complete this part only if this is a divisional, continuation or C-I-P application)

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) or PCT international application(s) designating the United States of America that is/are listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior application(s) in the manner provided by the first paragraph of Title 35, United States Code § 112, I acknowledge the duty to disclose information as defined in Title 37, Code of Federal Regulations, § 1.56 which occurred between the filing date of the prior application(s) and the national or PCT international filing date of this application:

(Application Serial No.) (Filing Date) (Status) (patented, pending, abandoned)

(Application Serial No.) (Filing Date) (Status) (patented, pending, abandoned)

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section

1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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